

**K952860 THE NINER NITINOL GUIDEWIRE**Dec 13, 1995  
174 days to decisionK952860 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k952860/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 22, 1995
Decision date	Dec 13, 1995
Days to decision	174 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Microvena Corp.</b>
Location	Findley, MN, US
Contact	JODI LOCHER
510(k) history	18 submissions · 18 cleared · 1990-1999

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952860/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 4, 2026